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PCT/AU 99/01033
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Re: Application of REINHARD, Judith; LACEY, Michael James and LENZ Michael
TERMITE ATTRACTANT AND/OR FEEDING STIMULANT
Assignee: COMMONWEALTH SCIENTIFIC AND INDUSTRIAL RESEARCH ORGANISATION
Our Ref: **Q65015**

Dear Sir:

The following documents and fees are submitted herewith in connection with the above application for the purpose of entering the National stage under 35 U.S.C. § 371 and in accordance with Chapter II of the Patent Cooperation Treaty:

X an English translation of the International Application (29 pages)

X an English translation of Article 34 amendments (annexes to the IPER).

X an International Search Report, Form PTO-1449 listing the ISR references.

It is assumed that copies of the International Application, the International Search Report, the International Preliminary Examination Report, and any Articles 19 and 34 amendments as required by § 371(c) will be supplied directly by the International Bureau, but if further copies are needed, the undersigned can easily provide them upon request.

Applicant claims benefit of small entity status in accordance with 37 CFR § 1.27.

The Government filing fee is calculated as follows **(Small Entity fees apply)**:

Total claims	47	-	20	=	27	x	$9.00	= $243.00
Independent claims	7	-	3	=	4	x	$40.00	= $160.00
Base Fee								$500.00
Multiple Dependent Claim Fee								$135.00
TOTAL FEE								$1038.00

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A check for the statutory filing fee of $1038.00 is attached. You are also directed and authorized to charge or credit any difference or overpayment to Deposit Account No. 19-4880. The Commissioner is hereby authorized to charge any fees under 37 C.F.R. §§ 1.16, 1.17 and 1.492 which may be required during the entire pendency of the application to Deposit Account No. 19-4880. A duplicate copy of this transmittal letter is attached.

Priority is claimed from December 22, 1998 based on Australian Application No. PP7842.

Respectfully submitted,

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TERMITE ATTRACTANT AND/OR FEEDING STIMULANT

TECHNICAL FIELD

The present invention is concerned with attractants and/or feeding stimulants for termites and, more particularly, with attractants and/or feeding stimulants for use in termite baits and as a component of termiticidal compositions.

BACKGROUND ART

Organochlorines have underpinned termite control around the world including in Australia, for many decades. With the ban on the use of organochlorines for termite control in Australia since 1995 and earlier or at similar times in other countries, increasing efforts are being mounted to develop alternative termite management systems. Bait systems for the control of active termite infestations are considered increasingly the key management option for such situations.

In bait systems termites are offered a matrix on which the insects ought to feed in preference to other food sources available to a termite colony. Termites either take up a slow-acting, non-repellent lethal product which is incorporated into the food (matrix) or the termites which aggregate in the matrix are directly treated with such a product. In both scenarios the agent is transported into the nest by the foragers and there distributed throughout the colony either via food exchange or mutual grooming between nest mates.

Following considerable research around the world there is now a growing awareness that just finding an effective bait toxin, initially thought to be the main impediment to the application of baits, is no guarantee at all that a

bait system will work effectively in practice. Control strategies relying on baits have to cope with the fact that termites have a choice and that the insects cannot be forced to make contact with the baits. Termites have to be able to locate a bait station in the first place, and once it is found, be attracted to it in significant numbers so that adequate transfer of the toxin from the bait site to the colony can occur. Differences in behaviour between species of termite, between colonies within a species and between conditions at various sites potentially restrict the effectiveness of this control strategy. Currently used bait matrices, in most cases just straight cellulose products (timber, cardboard, paper), do not necessarily ensure contact and build up of termite numbers in bait stations in a reliable, predictable fashion.

Attempts have been made to enhance the attraction of termites to bait matrices through the addition of attractant compounds. For example, International Application WO99/07218 describes the use of 2,4 heptadienal as an attractant for social pest insects such as wasps and termites. United States Patent No. 5,637,298 describes 2-4 naphthalenemethanol derivative substituted at the 7 or 8 position of the naphthalene ring structure by methyl, ethyl, propyl or isopropyl, and indicates that these compounds increase bait acceptance by termites. Likewise, United States Patent No. 5,756,114 describes the incorporation of certain aromatic compounds including resorcylic acid, protecatechuic acid and vanillic acid into baits on the basis that they act as food odour attractants. These compounds apparently mimic the trail-marking pheromone (*Z,Z,E*)-3,6,8-dodecatrien-1-ol. Thus, while they promote termite aggregation they do not necessarily stimulate feeding behaviour, and any increased feeding may

be a consequence only of the increased numbers of termites at a selected site.

Termites are social insects and the social organisation of termite colonies largely depends on chemical signals present in the environment or produced by members of the colony. These signals modulate a variety of behaviours including foraging for food or communal exploitation of a food source. For example, during feeding, termites release a chemical signal from an exocrine gland that stimulates nest mates to feed at the same site, thereby ensuring a rapid and efficient exploitation of the food source.

All species of termite have paired labial glands located in the thorax. The glandular ducts join in the head with those of the water sacs and the contents are secreted from the mouth as saliva. This secretion has been reported to have various functions depending on the species, and has variously been identified as a defensive substance in soldier termites, a regulator of nest microclimate, a supporter of fungal cultivation in the nest or as a social nutrient. In addition, the labial glands have been said to secrete a cementing substance for nest construction or gallery building and have been identified as a source of digestive enzymes.

More recently, Reinhard et al., *Journal of Chemical Ecology, Vol. 23 No. 10, 1997* concluded that the labial gland secretion may play a pheromonal role during food exploitation, and that this might be a general phenomenon in termites. Reinhard et al. took labial gland extracts and used these in feeding choice tests. They observed that the labial gland secretion carries a signal that stimulates gnawing and feeding by termite workers during food exploitation. The extract of the labial gland even

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elicited feeding behaviour when applied without food onto glass plates. These extracts were tested with both *Reticulitermes santonensis* and *Schedorhinotermes lamanianus* and proved to elicit a significant feeding preference in the two species. In view of this, Reinhard et al. suggested that the signal function of the labial gland secretion for food exploitation is phylogenetically old and non-species specific. The chemical signal has now been identified for the first time and has proved to work as a powerful feeding stimulant at natural low concentrations on a wide range of termite species. In view of this a class of compounds which stimulate termite feeding has been identified.

DISCLOSURE OF THE INVENTION

According to a first aspect of the present invention there is provided a feeding stimulant for stimulating feeding activity in termites, comprising an effective amount of a compound capable of stimulating feeding activity in termites, said compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, and a biologically acceptable carrier and/or extender.

Where the feeding stimulant is a compound in which at least one R is an organic group it may have feeding stimulating activity or may be a pre-cursor of a compound with feeding stimulating activity.

In the former case, the organic group is preferably selected from the group consisting of alkyl, substituted alkyl, aryl or substituted aryl, and in the latter case is typically a compound which is hydrolysed to one having feeding stimulating activity, such as those in which the

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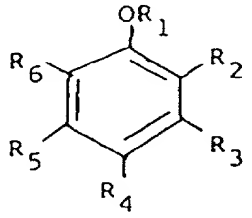
such compound. Polymers or oligomers such as polyphenylethers, as well as being long-lived in the

environment, will progressively hydrolyse to compounds having feeding stimulating activity.

Compounds having feeding stimulating activity typically comprise a benzene ring substituted by said at least two OR groups.

Typically such compounds have the following general formula I:



(I)

wherein R_1 is selected from the group consisting of hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, aralkyl, and substituted aralkyl;

R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, substituted alkyl, alkoxy, substituted alkoxy, aryl, substituted aryl, aryloxy, substituted aryloxy, aralkyl, substituted aralkyl, aralkyloxy and substituted aralkyloxy, or R_2 and R_3 together, R_3 and R_4 together, R_4 and R_5 together and/or R_5 and R_6 together form an aryl group;

provided only that at least one of R_2, R_3, R_4, R_5 or R_6 is hydroxyl, alkoxy, substituted alkoxy, aryloxy, substituted aryloxy, aralkyloxy or substituted aralkyloxy.

Preferably, R_1 is selected from the group consisting of hydrogen, alkyl, aryl and aralkyl.

More preferably, R_1 is selected from the group consisting of hydrogen, methyl, ethyl, phenyl and benzyl.

More preferably still, R_1 is hydrogen.

Preferably, R_2, R_3, R_4, R_5 and R_6 are independently

selected from the group consisting of hydrogen, hydroxyl, alkyl, alkoxy, aryl, aryloxy, aralkyl, and aralkyloxy.

More preferably, R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, methyl, ethyl, methoxy, ethoxy, phenyl, phenoxy, benzyl and benzyloxy.

More preferably still, at least one of R_2, R_3, R_4, R_5 or R_6 is hydroxyl. In particular, R_2 or R_6, R_3 or R_5 or R_4 is typically hydroxyl.

Particularly preferred compounds for use in the present invention are selected from the group consisting of:

p-hydroquinone (1,4-dihydroxybenzene)
catechol (1,2-dihydroxybenzene)
resorcinol (1,3-diahydroxybenzene)
phloroglucinol (1,3,5-trihydroxybenzene)
4-methoxyphenol
methoxyhydroquinone (1-methoxy-2,5-dihydroxybenzene)
1,4-dimethoxybenzene
4-phenoxyphenol
phenylhydroquinone
4-benzyloxyphenol

Moreover, addition compounds such as quinhydrone (an addition compound of 1 mole hydroquinone and 1 mole quinone) are also envisaged.

Alternatively, said compound may have a plurality of aryl moieties.

Preferably each said aryl moiety is a benzene ring and the compound is a polyphenylether. Typically, the polyphenylether is an ether of p-hydroquinone having between 2 and 5 p-hydroquinone residues.

As used throughout the specification and claims the term "alkyl" refers to straight or branched chain alkyl radicals, preferably C_1-C_{10} alkyl radicals and, more preferably, C_1-C_4 alkyl radicals.

As used throughout the specification and claims the term "substituted alkyl" refers to an alkyl radical substituted by any substituent, conveniently, by hydroxyl, alkoxy, carboxy, carboxyalkyl, carbamoyl, carbamido, amino, mono- or di- alkyl substituted amino, halogen, alkylcarbonyloxy or alkylcarbonylamino.

As used throughout the specification the term "aryl" refers to a six-membered carbocyclic aromatic ring or a five- or six-membered heterocyclic aromatic ring containing 1, 2 or 3 oxygen, nitrogen or sulphur atoms as the heteroatom, and includes fused ring systems containing a plurality of such rings.

As used throughout the specification and claims the term "substituted aryl" refers to an aryl radical substituted by any substituent, conveniently, by alkyl, hydroxyl, alkoxy, carboxy, carboxyalkyl, carbamoyl, carbamido, amino, mono- or di- alkyl substituted amino, halogen, alkylcarbonyloxy or alkylcarbonylamino.

As used throughout the specification and claims the term "alkoxy" refers to an alkoxy radical containing a straight or branched chain alkyl radicals, preferably C_1-C_{10} alkyl radicals and, more preferably, C_1-C_4 alkyl radicals.

As used throughout the specification and claims the term "substituted alkoxy" refers to an alkoxy radical substituted by any substituent, conveniently, by hydroxyl, alkoxy, carboxy, carboxyalkyl, carbamoyl, carbamido, amino, mono- or di- alkyl substituted amino, halogen, alkylcarbonyloxy or alkylcarbonylamino.

As used throughout the specification and claims the term "aryloxy" refers to an aryloxy radical containing a six-membered carbocyclic aromatic ring or a five- or six-membered heterocyclic aromatic ring containing 1, 2 or 3 oxygen, nitrogen or sulphur atoms as the heteroatom, and includes fused ring systems containing a plurality of such rings.

As used throughout the specification and claims the term "substituted aryloxy" refers to an aryloxy radical substituted by any substituent, conveniently, by alkyl, hydroxyl, alkoxy, carboxy, carboxyalkyl, carbamoyl, carbamido, amino, mono- or di- alkyl substituted amino, halogen, alkylcarbonyloxy or alkylcarbonylamino.

As used throughout the specification and claims the term "aralkyl" refers to an aralkyl radical comprising a straight or branched chain alkylene radical, preferably a C_1-C_{10} alkylene radical and, more preferably, a C_1-C_4 alkylene radical and a six-membered carbocyclic aromatic ring or a five- or six-membered heterocyclic aromatic ring containing 1, 2 or 3 oxygen, nitrogen or sulphur atoms as the heteroatom, and includes fused ring systems containing a plurality of such rings.

As used throughout the specification and claims the term "substituted aralkyl" refers to an aralkyl radical substituted by any substituent, conveniently, by hydroxyl, alkoxy, carboxy, carboxyalkyl, carbamoyl, carbamido, amino, mono- or di- alkyl substituted amino, halogen, alkylcarbonyloxy or alkylcarbonylamino.

As used throughout the specification and claims the term "aralkyloxy" refers to an aralkyloxy radical containing a straight or branched chain alkyleneoxy group, preferably a C_1-C_{10} alkyleneoxy group and, more preferably, C_1-C_4 alkyleneoxy group, and a six-membered carbocyclic

aromatic ring or a five- or six-membered heterocyclic aromatic ring containing 1, 2 or 3 oxygen, nitrogen or sulphur atoms as the heteroatom, and includes fused ring systems containing a plurality of such rings.

As used throughout the specification and claims the term "substituted aralkyloxy" refers to an aralkyloxy radical substituted by any substituent, conveniently, by hydroxyl, alkoxy, carboxy, carboxyalkyl, carbamoyl, carbamido, amino, mono- or di- alkyl substituted amino, halogen, alkylcarbonyloxy or alkylcarbonylamino.

As used throughout the specification and claims, the words "comprise", "comprises" and "comprising" are used in a non-exclusive sense, except where the context requires otherwise.

According to a second aspect of the present invention there is provided a method of stimulating feeding activity in termites, comprising the steps of:

(1) providing a feeding stimulant as described above; and

(2) applying said feeding stimulant to a locus.

Preferably, there is a food source at said locus.

According to a third aspect of the present invention there is provided a method of attracting termites to a locus, comprising the steps of:

(1) providing a food source at said locus,

(2) providing a feeding stimulant as described above; and

(3) applying said feeding stimulant to said locus.

The compounds of general formula I act as a feeding stimulant and/or attractant to termite species, in particular, to *Mastotermes darwiniensis*, *Coptotermes acinaciformis*, *Kalotermes flavicollis*, *Cryptotermes brevis*, *Hodotermes mossambicus*, *Zootermopsis angusticollis*,

Reticulitermes flavipes, *Reticulitermes santonensis*, *Heterotermes indicola*, *Schedorhinotermes lamanianus*, *Coptotermes formosanus*, *Nasutitermes nigriceps*, *Nasutitermes exitiosus*, *Trinervitermes trinervoides* and *Macrotermes subhyalinus*.

According to a fourth aspect of the present invention there is provided a bait for attracting termites, comprising:

(1) a food source; and

(2) a feeding stimulant as described above.

Typically the food source is a source of cellulose such as paper, cardboard, canite, chipboard, and sound or fungally decayed wood. The compound of general formula I is applied to the bait matrix in any convenient manner, such as by spraying a solution of the compound on the bait matrix, soaking the bait matrix in such a solution or by admixture with a solid compound of general formula I.

The bait matrix may also contain synergists and other attractants, as well as beneficial components such as nitrogen-containing compounds, carbohydrates and the like as nutrients.

Where necessary, antioxidants such as BHT, BHA or tocopherols may be added to stabilise the active compound within the bait. A controlled release system for the compound of general formula I may be employed where desirable.

Preferably, the bait matrix includes added toxins such as chitin synthesis inhibitors, insect growth regulators and other termiticides. Alternatively, termiticidal substances can be applied to the bait matrix once it has been deployed in the field and has attracted a significant number of termites. In either case, it is preferred that the toxin be slow-acting and non-repellent so as to be

transported into the nest by foragers and there distributed throughout the colony either via food exchange or mutual grooming between the nest mates.

According to a fifth aspect of the present invention there is provided a termiticidal composition comprising:

(1) a termiticidal substance; and

(2) a feeding stimulant as described above.

According to a sixth aspect of the present invention there is provided a compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, when used for stimulating feeding activity in termites.

According to a seventh aspect of the present invention there is provided a compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, when used in an amount effective to stimulate feeding activity in termites to attract termites to a locus.

According to an eighth aspect of the present invention there is provided the use of a compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, in stimulating feeding activity in termites.

According to a ninth aspect of the present invention there is provided the use in an amount effective to stimulate feeding activity in termites of a compound capable of stimulating feeding activity in termites, said compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, to attract

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termites to a locus.

According to a tenth aspect of the present invention there is provided the use in an amount effective to stimulate feeding activity in termites of a compound capable of stimulating feeding activity in termites in the manufacture of a bait for attracting termites, said compound having at least two OR groups, each of which is a substituent of

an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof.

According to an eleventh aspect of the present invention there is provided the use in an amount effective to stimulate feeding activity in termites of a compound capable of stimulating feeding activity in termites in the manufacture of a termiticidal composition, said compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof.

According to a twelfth aspect of the present invention there is provided a method of stimulating feeding activity in termites, comprising the steps of:

(1) providing a compound effective in stimulating feeding activity in termites having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof; and

(2) applying said compound to a locus.

Typically, the compound having at least two OR groups is a compound of general formula I as described above.

It has been found that para-hydroquinone is the natural feeding stimulant, but exists in the labial glands of termites almost entirely as its glucose conjugate, 4-hydroxyphenyl-ß-D-glucopyranoside, which is commonly called β-arbutin. β-Arbutin and glucose conjugates of the other compounds of general formula I may also be used in the invention described above. In particular, β-arbutin or glucose conjugates of the other compounds of formula I can be incorporated into a bait matrix and, through slow decay generating an active compound of general formula I, could act as a slow-release system.

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BEST MODE FOR CARRYING OUT THE INVENTION

Preferred embodiments of the invention will now be described, by way of example only, with reference to the following examples.

Example 1 - Use of Labial Glands Extracts as Termite Attractants

In order to prepare labial gland extracts, termites were killed and the paired labial glands were removed. The labial glands were disrupted by freezing them for 15 minutes at -20°C and extracted with 0.6 ml of water for 12

hours at room temperature. Then the extract was frozen at -20°C until used. The labial gland extracts prepared and tested are listed in Table I. Each extract was chemically analysed for the presence of para-hydroquinone, and it was found to be present in all. Selected extracts were used in a bioassay to establish feeding choice, as indicated in Table 1, below.

Table 1: Labial gland extracts prepared and tested

Termite species	No. of glands Extracted	Chemically Analysed	Bioassayed
Kalotermes flavicollis	40	+	
Cryptotermes brevis	70	+	+
Mastotermes darwiniensis	30	+	+
Hodotermes mossambicus	40	+	
Zootermopsis angusticollis	40	+	
Reticulitermes flavipes	70	+	+
Reticulitermes santonensis	70	+	+
Heterotermes indicola	120	+	
Schedorhinotermes lamanianus	60	+	
Coptotermes formosanus	70	+	+
Coptotermes acinaciformis	80	+	+
Nasutitermes nigriceps	60	+	
Nasutitermes exitiosus	70	+	+
Trinervitermes trinervoides	30	+	
Macrotermes subhyalinus	40	+	

The methodology employed in the choice tests was that used by Reinhard et al. *supra*. In these experiments the termites were housed in a suitable container with access via a silicone tube to a foraging arena. In each experiment two semicircles of moist filter paper (2.5cm in diameter) were placed close beside each other in the arena. One of the two semicircles was randomly chosen for application of one of the 25µl aliquots of labial gland extract and then

moistened with water. The other semicircle was just moistened. Feeding in termites is expressed by gnawing behaviour, which can be easily recognised by the hypognathous head positions wherein the termites bore their mandibles into the food and wriggle their heads trying to tear off little pieces, which they can then transport back to the nest.

The distribution of the first 20 gnawing/feeding termites on the semicircles was registered. For example, it was observed that 19 of 20 *Mastotermes darwiniensis* termites responded by gnawing and eating the filter paper treated with one equivalent of its labial gland secretion while only one termite responded to the control. Similarly, 18 of 20 *C. acinaciformis* termites responded by gnawing and eating the filter paper treated with 2.5 equivalents of its labial gland secretion while 2 responded to the control. A further important observation was that termites of selected species also responded strongly in the bioassay to labial gland secretion from an unrelated species. For instance, *C. acinaciformis* termites responded to a test paper treated with one equivalent of *M. darwiniensis* gland secretion while *M. darwiniensis* termites responded to a test paper treated with 2.5 equivalents of *C. acinaciformis* gland secretion. These results demonstrate that the labial gland extract is a non-specific feeding stimulant for termites. The results are summarised in Table 2.

Table 2: Natural lures

Termite species Responding to lure	Origin of labial gland Extract	Quantity of extract (gland equivalents)	Response
M.darwiniensis	*M.darwiniensis*	1	+++
	C.acinaciformis	2.5	+++
C.acinaciformis	*M.darwiniensis*	1	+++
	C.acinaciformis	2.5	+++

An analysis of the labial gland extract shows that para-hydroquinone is present at low levels, usually less than 10^{-10} grams per gland, but is present at much higher concentrations in the saliva. β-arbutin is present in high concentrations in the glands but is no longer evident in the saliva. Presumably β-arbutin is broken down enzymatically into para-hydroquinone and glucose during release of the termite's saliva, hence it was postulated that para-hydroquinone was the principal chemical feeding stimulant.

Example 2 - Synthetic Compounds as Termite Attractants

Feeding choice tests were conducted with para-hydroquinone and a number of related chemical substances in the manner described above in Example 1. The experimental data is summarised in Table 3.

Table 3: Synthetic lures

Termite species responding to lure	Compound	Quantity in Lure[ng]	Response
M.darwiniensis	p-hydroquinone	5	+++
	Quinhydrone	5	++
	Catechol	5	-/+
	Resorcinol	5	+
	Phloroglucinol	5	-/+
	4-methoxyphenol	5	+
	Methoxyhydroquinone	5	+
	1,4-dimethoxybenzene	5	++
	4-phenoxyphenol	5	-/+
	phenylhydroquinone	5	+
	polyphenylether*	5	+
	4-benzyloxyphenol	5	-/+
C.acinaciformis	p-hydroquinone	1	+++
	quinhydrone	5	++
	catechol	5	++
	resorcinol	5	++
	phloroglucinol	5	++
	4-methoxyphenol	5	+
	methoxyhydroquinone	5	+
	1,4-dimethoxybenzene	5	+
	4-phenoxyphenol	5	-/+
	phenylhydroquinone	5	-
	polyphenylether*	5	-
	4-benzyloxyphenol	5	-
S.actuosus	p-hydroquinone	5	+++
C.brevis	p-hydroquinone	5	+++
N.exitiosus	p-hydroquinone	5	+++
R.santonensis	p-hydroquinone	5	+++
R.flavipes	p-hydroquinone	5	+++
C.formosanus	p-hydroquinone	5	+++

* Mixture comprising mainly a pentamer of p-hydroquinone, but including dimer and trimer of p-hydroquinone as impurities.

When synthetic lures were tested, none of the principal labial gland constituents (glucose, inositols, β-arbutin) elicited any feeding stimulation, except at unnaturally high concentrations where they probably served a nutritional role as food supplements. However p-hydroquinone elicited feeding stimulation at natural trace levels in the laboratory bioassays. For instance the threshold for attraction was 5 nanograms p-hydroquinone (50 picomoles) for *M. darwiniensis* and 100 picograms p-hydroquinone (1 picomole) for *C. acinaciformis*. Thus, there are different lower thresholds of feeding stimulation for different termite species.

Synthetic compounds somewhat related in molecular structure to hydroquinone also elicited feeding responses from *M. darwiniensis* and *C. acinaciformis* in the laboratory bioassays, as shown in Table 3.

Example 3 - Mode of Attraction

The mode of attraction of termites to the para-hydroquinone source may well include both olfactory and gustatory stimulation. The attractivity of para-hydroquinone over distance (olfactory perception) was tested both in empty and sand-filled plastic arenas (ID 14.5 cm, height 1 cm, covered with a glass plate), which were attached via a silicone tube to the housing container of the termites. Tests were carried out with *M. darwiniensis* and *C. acinaciformis*. Per test, two treated filter papers (25ng - 25µg p-hydroquinone and water as control, respectively) were placed in opposite positions in the arenas. The direction of the tunnel/galleries built and the behaviour of a foraging termites in reference to the position of the filter papers were evaluated. In all tests both termite species built tunnels/galleries in direction

to the p-hydroquinone-treated filter paper, never towards the control filter paper. When foraging the termites usually walked slowly in a zigzag way, but when in proximity of the source of p-hydroquinone (ca. 5-6 cm), their behaviour changed suddenly: they walked straight and fast to the treated filter paper. Based on these observational data we concluded that the vapour of p-hydroquinone creates an "active space" of several centimetres, which once perceived directs the termites towards the source of the vapour by the concentration gradient. This active space did not get larger with increased p-hydroquinone concentration.

Example 4 - Choice Feeding Tests

Laboratory colonies of *Mastotermes darwiniensis* and *Coptotermes acinaciformis* have been tested in a choice feeding test (mimicking an actual bait situation in the field) with pieces of *Eucalyptus regnans* wood (ca. 3.5g). The colonies (ca. 500 termites in *M. darwiniensis*, 2000 termites in *C. acinaciformis)* were housed in plastic containers. Plastic arenas of 5cm diameter, 3.5cm high were attached with perspex tubes on opposite sides of the colony container. In these arenas the wood was offered: one treated with 20ng p-hydroquinone, dissolved in water, the other just moistened as control. The wood was dried and weighed before and after the test, the difference in weight as the amount eaten by termites was analysed after 3 days, 1 week and 4 weeks.

After 3 days and one week both *M. darwiniensis* and *C. acinaciformis* had eaten significantly more of the wood treated with the feeding stimulant than of the control (See Table 4). After 4 weeks the effect was gone. Therefore p-hydroquinone does act as feeding stimulant in a choice

feeding test, although as only a little p-hydroquinone was applied, the effect was only short-term. This could be improved when testing the signal in the field under natural conditions and with complete termite colonies.

Table 4: Laboratory choice feeding tests with *Mastotermes darwiniensis* and *Coptotermes acinaciformis*: Amount wood eaten [g] after 3 days, one week and four weeks, comparing wood treated with 20ng p-hydroquinone to control (mean ± sd, n=20, Wilcoxon-Matched-Pairs-Test, ***: significant difference at $p<0.001$, n.s.: no significant difference).

Species	Duration of trial	Treated wood eaten [g]	Control wood eaten [g]	P
M.darwiniensis	3 days	0.234±0.139	0.121±0.108	***
	1 week	0.737±0.557	0.506±0.527	**
	4 weeks	2.397±0.968	2.255±0.918	n.s.
C.acinaciformis	3 days	0.056±0.0.36	0.032±0.033	**
	1 week	0.185±0.159	0.096±0.109	**
	4 weeks	1.162±0.851	1.209±0.929	n.s.

Example 5 - Field Trials

Colonies of *Coptotermes lacteus* (ACT), *Coptotermes acinaciformis* (NT) and *Mastotermes darwiniensis* (NT) have been used for large baiting trials in the field. Furthermore *Coptotermes frenchi* (ACT), *Nasutitermes exitiosus* (NSW), *Schedorhinotermes actuosus* (NT), *Coptotermes travians* (Malaysia) and *Coptotermes curvignathus* (Malaysia) have been tested exemplarily at infestation sites in urban areas and in the field. Paper towel of ca. 10g was used as bait matrix. It was either treated with 20µg hydroquinone (dissolved in water) or moistened with water only (=control). The paper was folded and stuffed in plastic tubes. Termites had access to the bait material through holes drilled into the tubes. One

treated and one control bait each were placed at feeding/infestation sites of the field colonies. In case of the larger field trials, up to 24 colonies per species had been selected, and drums filled with wood had been dug into the soil around colonies as feeding sites. Baits were placed on top of the infested drums and covered with plastic foil and soil. In case of the exemplary trials single infestation sites have been selected and the baits were attached directly onto the infestation and covered with plastic foil and soil, or cardboard to ensure minimum disturbance. Baits were checked after 1 to 4 days or after 2 weeks, depending on species and activity. The amount of paper eaten and the number of termites were analysed.

As usual in natural field colonies there was a strong variation of data between colonies, therefore for statistical analysis data had to be transformed into log and square root, respectively. In the large field trials *C. lacteus, M. darwiniensis and C. acinaciformis* had all consumed significantly more of the bait material and there were more termites attracted to the baits, when hydroquinone had been applied (See Table 5). The exemplary tests with *C. frenchi, S. actuosus, C. travians* and *C. curvignathus* all indicated increased feeding activity on treated baits over control baits (See Table 6). Tests with *N. exitiosus* showed no feeding activity even after long exposure, due to the difficult dietary preferences of this species. However, we could still show increased termite presence in treated baits compared to control baits (See Table 6). We therefore conclude that hydroquinone in fact acts also under natural conditions in the field as strong and effective attractant and feeding stimulant on various termite species, when added to baits.

Table 5: Field baiting trials with *Coptotermes lacteus* (ACT), *Coptotermes acinaciformis* (NT) and *Mastotermes darwiniensis* (NT). **(W)**: Amount bait material eaten [g] and **(No)**: number of termites [N} in baits, comparing baits treated with 20µg hydroquinone to control baits (mean ± SE, Paired Samples T-test, ***: significant difference at $p<0.001$, data transformed to log or sqrt for statistical analysis).

Species	**Trial**	***N***		**Treated Bait**	**Control Bait**	***P***
C.lacteus	3 days	17	**W[g]**	0.392±0.156	0.181±0.099	***
			No[N]	269.7±169.1	198.9±147.7	***
M.darwiniensis	2 days	12	**W[g]**	0.766±0.164	0.368±0.126	**
			No[N]	18.5±3.6	7.7±2.9	**
C.acinaciformis	2 days	16	**W[g]**	0.056±0.010	0.035±008	**
			No[N]	120.6±29.5	84.1±20.0	***

Table 6: Exemplary field baiting trials with *Coptotermes frenchi* (ACT), *Schedorhinotermes actuosus* (NT), *Coptotermes travians* (Malaysia), *Coptotermes curvignathus* (Malaysia) and *Nasutitermes exitiosus* (NSW). Proportion bait material eaten [%] or termite presence, respectively, comparing baits treated with 20µg hydroquinone to control baits.

Species	Trial	*n*	Proportion eaten/termite presence (treated bait)	Proportion eaten/termite presence (control bait)
C. frenchi	2 weeks	3	30%	5%
			90%	0%
			30%	0%
S. actuosus	4 days	1	5%	0%
C. travians	1 day	2	20%	20%
			60%	40%
C. curvignathus	1 day	3	95%	20%
			50%	5%
N. exitiosus	2 weeks	4	Termites present	Not touched
			Termites present	Not touched
			Termites present	Not touched
			Termites present	Not touched

INDUSTRIAL APPLICABILITY

The compounds of the present invention are useful in stimulating feeding activity in termites so as to enhance the effectiveness of termite baits.

ARTICLE 34

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A feeding stimulant for stimulating feeding activity in termites, comprising an effective amount of a compound capable of stimulating feeding activity in termites, said compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, and a biologically acceptable carrier and/or extender.

2. A feeding stimulant as claimed in claim 1 wherein at least one R is an organic group and said compound has feeding stimulating activity.

3. A feeding stimulant as claimed in claim 2 wherein said organic group is selected from the group consisting of alkyl, substituted alkyl, aryl, substituted aryl, aralkyl and substituted aralkyl.

4. A feeding stimulant as claimed in claim 1 wherein at least one R is an organic group and said compound is a precursor of a compound with feeding stimulating activity.

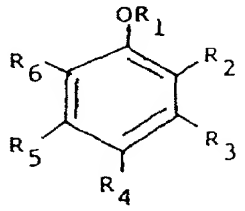
5. A feeding stimulant as claimed in claim 4 wherein said compound is hydrolysed to a compound in which said at least one R is hydrogen.

6. A feeding stimulant as claimed in claim 5 wherein said organic group is a carbohydrate moiety.

7. A feeding stimulant as claimed in claim 6 wherein said compound is β-arbutin.

8. A feeding stimulant as claimed in claim 1 wherein said aryl group is a benzene ring substituted by said at least two OR groups.

9. A feeding stimulant as claimed in claim 8 wherein said compound has the following general formula I:



(I)

wherein R_1 is selected from the group consisting of hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, aralkyl and substituted aralkyl;

R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, substituted alkyl, alkoxy, substituted alkoxy, aryl, substituted aryl, aryloxy, substituted aryloxy, aralkyl, substituted aralkyl, aralkyloxy and substituted aralkyloxy, or R_2 and R_3 together, R_3 and R_4 together, R_4 and R_5 together and/or R_5 and R_6 together form an aryl group;

provided only that least one of R_2, R_3, R_4, R_5 or R_6 is hydroxyl, alkoxy, substituted alkoxy, aryloxy, substituted aryloxy, aralkyloxy or substituted aralkyloxy.

10. A feeding stimulant as claimed in claim 9 wherein R_1 is selected from the group consisting of hydrogen, alkyl, aryl and aralkyl.

11. A feeding stimulant as claimed in claim 10 wherein R_1 is selected from the group consisting of hydrogen, methyl, ethyl, phenyl and benzyl.

12. A feeding stimulant as claimed in claim 11 wherein R_1 is hydrogen.

13. A feeding stimulant as claimed in claim 9 wherein R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, alkoxy, aryl, aryloxy, aralkyl, and aralkyloxy.

14. A feeding stimulant as claimed in claim 13 wherein R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, methyl, ethyl, methoxy, ethoxy, phenyl, phenoxy, benzyl and benzyloxy.

15. A feeding stimulant as claimed in claim 14 wherein R_2 or R_6 is hydroxyl.

16. A feeding stimulant as claimed in claim 14 wherein R_3 or R_5 is hydroxyl.

17. A feeding stimulant as claimed in claim 14 wherein R_4 is hydroxyl.

18. A feeding stimulant as claimed in claim 1 wherein said compound is selected from the group consisting of:

p-hydroquinone
quinhydrone
catechol
resorcinol
phloroglucinol
4-methoxyphenol
methoxyhydroquinone
1,4-dimethoxybenzene

4-phenoxyphenol
phenylhydroquinone
4-benzyloxyphenol

19. A feeding stimulant as claimed in claim 1 wherein said compound has a plurality of aryl moieties.

20. A feeding stimulant as claimed in claimed 19 wherein each said aryl moiety is a benzene ring.

21. A feeding stimulant as claimed in claim 20 wherein said compound is a polyphenylether.

22. A method of stimulating feeding activity in termites, comprising the steps of:

(1) providing a feeding stimulant as claimed in any one of claims 1 to 21; and

(2) applying said feeding stimulant to a locus.

23. A method as claimed in claim 22 further comprising the step of providing a food source at said locus.

24. A method of attracting termites to a locus, comprising the steps of:

(1) providing a food source at said locus;

(2) providing a feeding stimulant as claimed in any one of claims 1 to 21; and

(3) applying said feeding stimulant to said locus.

25. A bait for attracting termites, comprising:

(1) a food source; and

(2) a feeding stimulant as claimed in any one of claims 1 to 21.

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claims 1 to 21.

26. A bait as claimed in claim 25 wherein said food source is a source of cellulose.

27. A bait as claimed in claim 26 wherein said food source is selected from the group consisting of paper, cardboard, canite, chipboard, sound wood and fungally decayed wood.

28. A bait as claimed in any one of claims 25 to 27 further comprising a termiticidal substance.

29. A bait as claimed in claim 28 in which said termiticidal substance is a chitin synthesis inhibitor or an insect growth regulator.

30. A bait as claimed in any one of claims 25 to 29 further comprising an antioxidant.

31. A bait as claimed in any one of claims 25 to 30 further comprising a synergist and/or other attractants.

32. A bait as claimed in any one of claims 25 to 31 further comprising nutrients such as nitrogen-containing compounds and carbohydrates.

33. A termiticidal composition comprising:

(1) a termiticidal substance; and

(2) a feeding stimulant as claimed in any one of claims 1 to 21.

34. A termiticidal composition as claimed in claim 33 wherein said termiticidal substance is a chitin synthesis inhibitor or insect growth regulator.

35. A compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, when used for stimulating feeding activity in termites.

36. A compound as claimed in claim 35 of general formula I as defined in claim 9.

37. A compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, when used in an amount effective to stimulate feeding activity in termites to attract termites to a locus.

38. A compound as claimed in claim 37 of general formula I as defined in claim 9.

39. The use of a compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, in stimulating feeding activity in termites.

40. The use of a compound as claimed in claim 39 wherein said compound is of general formula I as defined in claim 9.

41. The use in an amount effective to stimulate feeding activity in termites of a compound capable of stimulating feeding activity in termites, said compound having at least

two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof, to attract termites to a locus.

42. The use of a compound as claimed in claim 41 wherein said compound is of general formula I as defined in claim 9.

43. The use in an amount effective to stimulate feeding activity in termites of a compound capable of stimulating feeding activity in termites in the manufacture of a bait for attracting termites, said compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof.

44. The use of compound as claimed in claim 43 wherein said compound is of general formula I as defined in claim 9.

45. The use in an amount effective to stimulate feeding activity in termites of a compound capable of stimulating feeding activity in termites in the manufacture of a termiticidal composition, said compound having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof.

46. The use of a compound as claimed in claim 45 wherein said compound is of general formula I as defined in claim 9.

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47. A method of stimulating feeding activity in termites, comprising the steps of:

(1) providing a compound effective in stimulating feeding activity in termites having at least two OR groups, each of which is a substituent of an aryl moiety, and R is hydrogen or an organic group, and addition compounds thereof; and

(2) applying said compound to a locus.

48. A method as claimed in claim 47 wherein at least one R is an organic group and said compound has feeding stimulating activity.

49. A method as claimed in claim 48 wherein said organic group is selected from the group consisting of alkyl, substituted alkyl, aryl, substituted aryl, aralkyl and substituted aralkyl.

50. A method as claimed in claim 47 wherein at least one R is an organic group and said compound is a precursor of a compound with feeding stimulating activity.

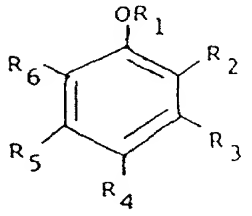
51. A method as claimed in claim 50 wherein said compound is hydrolysed to a compound in which said at least one R is hydrogen.

52. A method as claimed in claim 51 wherein said organic group is a carbohydrate moiety.

53. A method as claimed in claim 52 wherein said compound is β-arbutin.

54. A method as claimed in claim 47 wherein said compound has an aromatic nucleus substituted by said at least [illegible] groups.

55. A method as claimed in claim 54 wherein said compound has the following general formula I:



(I)

wherein R_1 is selected from the group consisting of hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, aralkyl and substituted aralkyl;

R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, substituted alkyl, alkoxy, substituted alkoxy, aryl, substituted aryl, aryloxy, substituted aryloxy, alkaryl, substituted alkaryl, alkaryloxy and substituted alkaryloxy, or R_2 and R_3 together, R_3 and R_4 together, R_4 and R_5 together and/or R_5 and R_6 together form an aryl group;

provided only that least one of R_2, R_3, R_4, R_5 or R_6 is hydroxyl, alkoxy, substituted alkoxy, aryloxy, substituted aryloxy, alkaryloxy or substituted alkaryloxy.

56. A method as claimed in claim 55 wherein R_1 is selected from the group consisting of hydrogen, alkyl, aryl and alkaryl.

57. A method as claimed in claim 56 wherein R_1 is selected from the group consisting of hydrogen, methyl, ethyl,

phenyl and benzyl.

58. A method as claimed in claim 57 wherein R_1 is hydrogen.

59. A method as claimed in claim 54 wherein R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, alkyl, alkoxy, aryl, aryloxy, alkaryl, and alkaryloxy.

60. A method as claimed in claim 58 wherein R_2, R_3, R_4, R_5 and R_6 are independently selected from the group consisting of hydrogen, hydroxyl, methyl, ethyl, methoxy, ethoxy, phenyl, phenoxy, benzyl and benzyloxy.

61. A method as claimed in claim 60 wherein R_2 or R_6 is hydroxyl.

62. A method as claimed in claim 60 wherein R_3 or R_5 is hydroxyl.

63. A method as claimed in claim 60 wherein R_4 is hydroxyl.

64. A method as claimed in claim 47 wherein said compound is selected from the group consisting of:

p-hydroquinone
quinhydrone
catechol
resorcinol
phloroglucinol
4-methoxyphenol
methoxyhydroquinone
1,4-dimethoxybenzene

4-phenoxyphenol
phenylhydroquinone
4-benzyloxyphenol

65. A method as claimed in claim 47 wherein said compound has a plurality of aryl moieties.

66. A method as claimed in claimed 42 wherein each said aryl moiety is a benzene ring.

67. A method as claimed in claim 43 wherein said compound is a polyphenylether.

SOLE/JOINT

DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I hereby declare that my residence, mailing address and citizenship are as stated below next to my name: that I verily believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter claimed and for which a patent is sought in the application entitled:

TERMITE ATTRACTANT AND/OR FEEDING STIMULANT

which application is:

☐ the attached application
(for original application)

Application No. International PCT/AU/01033
(Confirmation No. Not assigned) filed November 25, 1999, and amended on

(for declaration not accompanying application)

that I have reviewed and understand the contents of the specification of the above-identified application, including the claims, as amended by any amendment referred to above; that I acknowledge my duty to disclose information of which I am aware and which is material to the patentability of this application as defined in 37 C.F.R. 1.56, that I hereby claim priority benefits under Title 35, United States Code §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate, §119(e) of any United States provisional application(s), or §365(a) of any PCT International application which designated at least one country other than the United States of America, listed below and have also identified below any foreign application for patent or inventor's certificate or of any PCT International application having a filing date before that of the application on which priority is claimed:

Application Number	Country	Filing Date	Priority Claimed Yes	Priority Claimed No
PP7842	AU	December 22, 1998	X	☐

I hereby claim the benefit under 35 United States Code §120 of any United States application(s), or §365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in a listed prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge my duty to disclose any information material to the patentability of this application as defined in 37 C.F.R. 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

Application No.	Filing Date	Status

33

I hereby appoint John H. Mion, Reg. No. 18,879; Thomas J. Macpeak, Reg. No. 19,292; Robert J. Seas, Jr., Reg. No. 21,092; Darryl Mexic, Reg. No. 23,063; Robert V. Sloan, Reg. No. 22,775; Peter D. Olexy, Reg. No. 24,513; J. Frank Osha, Reg. No. 24,625; Waddell A. Biggart, Reg. No. 24,861; Louis Gubinsky, Reg. No. 24,835; Neil B. Siegel, Reg. No. 25,200; David J. Cushing, Reg. No. 28,703; John R. Inge, Reg. No. 26,916; Joseph J. Ruch, Jr., Reg. No. 26,577; Sheldon I. Landsman, Reg. No. 25,430; Richard C. Turner, Reg. No. 29,710; Howard L. Bernstein, Reg. No. 25,665; Alan J. Kasper, Reg. No. 25,426; Kenneth J. Burchfiel, Reg. No. 31,333; Gordon Kit, Reg. No. 30,764; Susan J. Mack, Reg. No. 30,951; Frank L. Bernstein, Reg. No. 31,484; Mark Boland, Reg. No. 32,197; William H. Mandir, Reg. No. 32,156; Brian W. Hannon, Reg. No. 32,778; Abraham J. Rosner, Reg. No. 33,276; Bruce E. Kramer, Reg. No. 33,725; Paul F. Neils, Reg. No. 33,102; Brett S. Sylvester, Reg. No. 32,765; Robert M. Masters, Reg. No. 35,603, George F. Lehnigk, Reg. No. 36,359, John T. Callahan, Reg. No. 32,607, Steven M. Gruskin, Reg. No. 36,818, Peter A. McKenna, Reg. No. 38,551 and Edward F. Kenehan, Reg. No. 28,962, my attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith, and request that all correspondence about the application be addressed to **SUGHRUE, MION, ZINN, MACPEAK & SEAS, PLLC**, 2100 Pennsylvania Avenue, N.W., Washington, D.C. 20037-3213.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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